

GenCore version 5.1.4-p5.4578
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OM protein - protein search, using sw model

Run on: April 8, 2003, 14:25:27 ; Search time 75 Seconds

(without alignments)
968,288 Million cell updates/sec

Title: US-09-001-737-8

Perfect score: 545

Sequence: 1 MAKEIFSADARAAVAVRGVD.....TPAPAMPAGMDPGMGMGNC 545

Scoring table: OLIGO

Gapop 60.0 , Capext 60.0

Searched: 908470 seqs, 133250620 residues

Word size: 8

Total number of hits satisfying chosen parameters: 350

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 1000 summaries

Database:

A_Geneseq-101002:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	545	100.0	545	20	AAV23904
2	538	98.7	545	23	ABP28529
3	78	14.3	540	23	ABP28528
4	70	12.8	540	22	AAAM01101
5	70	12.8	541	20	AAV23902
6	70	12.8	641	22	ABR31619
7	56	10.3	542	23	ABR53701
8	31	5.7	542	23	ABR49241
9	26	4.8	539	20	AAV23916
10	24	4.4	224	20	AAV37100

11	24	4.4	419	18	AAV10977	Dihydrofolate redu
12	24	4.4	419	18	AAV11865	DHFR/Polypeptide B
13	24	4.4	419	12	AAV13337	Hypr protein Chl
14	24	4.4	544	16	AAV67383	C. psittaci Hypr g
15	24	4.4	544	18	AAV10975	Chlamydia pneumoni
16	24	4.4	544	18	AAV11865	Polypeptide B. Ch
17	24	4.4	544	20	AAV23905	Amino acid sequenc
18	24	4.4	544	22	AAE11757	Amino acid sequenc
19	24	4.4	544	23	ABR94272	Chlamydia pneumoni
20	24	4.4	545	21	AAV19080	Amino acid sequenc
21	24	4.4	704	18	AAV10976	Amino acid sequenc
22	24	4.4	704	18	AAV11864	Dihydrofolate redu
23	24	4.2	547	22	AAV65060	DHFR/Polypeptide B
24	23	4.2	574	22	AAV65061	Pseudomonas aerugi
25	23	4.2	612	22	AAV63908	ptichisB expressio
26	23	3.9	301	22	AAV63904	S. epidermidis ope
27	21	3.9	539	22	AAV61848	S. epidermidis ope
28	21	3.9	541	20	AAV23917	Amino acid sequenc
29	21	3.9	545	20	AAV23930	Amino acid sequenc
30	21	3.7	309	19	AAV60146	Consensus mlna act
31	20	3.7	309	19	AAV14893	M. vaccae antigen
32	20	3.7	309	20	AAV73499	Amino acid sequenc
33	20	3.7	327	20	AAV14910	M. vaccae GroEL hom
34	20	3.7	327	20	AAV73516	Amino acid sequenc
35	20	3.7	523	19	AAV60144	M. vaccae GroEL hom
36	20	3.7	523	20	AAV14891	M. vaccae antigen
37	20	3.7	523	23	AAV73497	Amino acid sequenc
38	20	3.7	539	20	AAV23919	M. vaccae GroEL hom
39	20	3.7	540	9	AAV81351	Amino acid sequenc
40	20	3.7	540	16	AAV81610	Sequence of Mycob
41	20	3.7	540	16	AAV82100	Sequence of Mycob
42	20	3.7	540	19	AAV44702	Mycobacterium tube
43	20	3.7	540	18	AAV44702	Mycobacterium tube
44	20	3.7	540	21	AAV23932	Amino acid sequenc
45	20	3.7	540	21	AAV93332	Amino acid sequenc
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83	20	3.7	540	22	AAE11755	Myobacterium tube

84	17	3.1	548	17	AAB94368	Brevibacterium fla
85	17	3.1	548	22	AA92732	C. glutamicum prote
86	17	3.1	549	21	AA99642	B. pseudomallei Gr
87	16	2.9	16	17	AA94778	Peptide from libra
88	16	2.9	16	18	AAW43456	Mycobacteria sp. h
89	16	2.9	19	17	AA94827	Peptide from libra
90	16	2.9	19	18	AAW43505	Mycobacteria sp. h
91	16	2.9	51	22	AAB3742	Chaperone cpn60 pr
92	16	2.9	52	13	AAB70195	Heat shock protein
93	16	2.9	103	21	AAB41321	Human ORFX ORF1085
94	16	2.9	103	23	ABP03730	Human ORFX protein
95	16	2.9	112	19	AAW60170	M. vaccae GroEL-ho
96	16	2.9	112	20	AAW44876	M. vaccae GroEL-ho
97	16	2.9	112	23	ABW34482	M. vaccae GroEL-ho
98	16	2.9	118	23	ABP06879	Human ORFX protein
99	16	2.9	215	19	AAW60145	M. vaccae GroEL-ho
100	16	2.9	215	20	AAW44892	M. vaccae GroEL-ho
101	16	2.9	215	23	ABW34488	M. vaccae GroEL-ho
102	16	2.9	295	22	AAB31615	Amino acid sequenc
103	16	2.9	440	13	AAW23362	Amino acid sequenc
104	16	2.9	539	20	AAW23362	Amino acid sequenc
105	16	2.9	540	13	AAW23362	Amino acid sequenc
106	16	2.9	544	21	AAW75745	Neisseria meningit
107	16	2.9	544	21	AAW75747	Neisseria meningit
108	16	2.9	545	20	AAW233915	Amino acid sequenc
109	16	2.9	548	18	AAW16678	Amino acid sequenc
110	16	2.9	15	17	AAW97488	Lawsonia intracell
111	16	2.9	15	23	AAW99970	Mycobacterium heat
112	16	2.9	19	17	AAW44871	Peptide from libra
113	16	2.9	19	18	AAW43549	Mycobacteria sp. h
114	16	2.9	20	18	AAW43549	Mycobacteria sp. h
115	16	2.9	20	18	AAW43549	Mycobacteria sp. h
116	16	2.9	152	23	ABW42335	Human hsp60 peptid
117	16	2.9	160	19	AAW41375	Human ovarian anti
118	16	2.9	160	19	AAW65071	GroEL apical domai
119	16	2.9	186	19	AAW61377	E. coli GroEL prot
120	16	2.9	186	19	AAW65069	GroEL apical regio
121	16	2.9	544	20	AAW23903	E. coli GroEL N-te
122	16	2.9	545	20	AAW23903	Streptococcus pyog
123	16	2.9	547	16	AAW67381	Streptococcus pneu
124	16	2.9	547	16	AAW67381	L. pneumophila Hsp
125	16	2.9	547	16	AAW67381	Mitochondrial prot
126	16	2.9	548	23	AAW76192	Human P1 protein (
127	16	2.9	548	23	AAW76192	E. coli GroEL gene
128	16	2.9	548	19	AAW61378	E. coli GroEL prot
129	16	2.9	548	19	AAW65070	E. coli GroEL prot
130	16	2.9	548	20	AAW23908	Amino acid sequenc
131	16	2.9	548	20	AAW23908	MO9902989 Seg ID 1
132	16	2.9	548	21	AAW96621	Escherichia coli g
133	16	2.9	549	22	AAW50536	Escherichia coli g
134	16	2.9	559	20	AAW23909	Amino acid sequenc
135	16	2.9	559	16	AAW64764	E. coli GroEL prot
136	16	2.9	573	11	AAW44713	Amino acid sequenc
137	16	2.9	573	18	AAW14946	Human heat shock p
138	16	2.9	573	18	AAW12345	Human heat shock p
139	16	2.9	573	18	AAW12345	Human heat shock p
140	16	2.9	573	19	AAW56120	Human heat shock p
141	16	2.9	573	20	AAW23926	Protein sequenc o
142	16	2.9	573	21	AAW93333	Amino acid sequenc
143	16	2.9	573	21	AAW93333	Amino acid sequenc
144	16	2.9	573	22	AAW58685	Drosophila melanog
145	16	2.9	573	22	AAW58685	Drosophila melanog
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150	16	2.9	573	22	AAW58685	Drosophila melanog
151	16	2.9	573	22	AAW58685	Drosophila melanog
152	16	2.9	573	22	AAW58685	Drosophila melanog
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156	16	2.9	573	22	AAW58685	Drosophila melanog

230	11	2.0	19	22	AAH8291	Hsp-65 peptide epi
231	11	2.0	20	16	AAH8161	Heat shock protein
232	11	2.0	22	21	AAI93328	Amino acid sequenc
233	11	2.0	65	20	AAH89955	Expressed antigen
234	11	2.0	99	21	AAI12294	Zea mays protein f
235	11	2.0	104	21	AAI12293	Zea mays protein f
236	11	2.0	114	11	AAH07688	Mycobacterial 65KD
237	11	2.0	114	11	AAH07689	Mycobacterial 65KD
238	11	2.0	114	11	AAH07690	Mycobacterial 65KD
239	11	2.0	118	22	AAU02875	Synthetic autoanti
240	11	2.0	120	23	AAH14677	M. vaccae 65Kd he
241	11	2.0	120	23	AAH73483	Human cancer assoc
242	11	2.0	135	21	AAH44098	Human cancer assoc
243	11	2.0	156	21	AAH44097	Zea mays protein f
244	11	2.0	177	21	AAH08699	Human secreted pro
245	11	2.0	212	22	AAH63522	Human gastric can
246	11	2.0	247	22	AAH63519	Human gastric can
247	11	2.0	252	22	AAH14357	Novel human diagn
248	11	2.0	258	20	AAH37099	Amino acid sequenc
249	11	2.0	261	22	AAU02079	Synthetic multi-ta
250	11	2.0	342	22	AAH14356	Novel human diagn
251	11	2.0	397	21	AAH53326	Arabidopsis thalia
252	11	2.0	412	21	AAH49547	Arabidopsis thalia
253	11	2.0	423	21	AAH53325	Arabidopsis thalia
254	11	2.0	443	21	AAH49136	Arabidopsis thalia
255	11	2.0	446	21	AAH50252	Arabidopsis thalia
256	11	2.0	459	22	AAU02077	Synthetic multi-ta
257	11	2.0	464	22	AAU45673	Protonlbacterium
258	11	2.0	469	21	AAH49135	Arabidopsis thalia
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260	11	2.0	536	21	AAH50259	Arabidopsis thalia
261	11	2.0	536	21	AAH49134	Arabidopsis thalia
262	11	2.0	541	21	AAH50251	Arabidopsis thalia
263	11	2.0	568	20	AAH23924	Amino acid sequenc
264	11	2.0	586	21	AAH19771	Arabidopsis thalia
265	11	2.0	586	21	AAH50258	Arabidopsis thalia
266	11	2.0	611	21	AAH19770	Arabidopsis thalia
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270	11	2.0	650	21	AAH50250	Arabidopsis thalia
271	11	2.0	650	21	AAH50250	Arabidopsis thalia
272	11	1.8	10	22	AAH88269	Hsp-65 peptide epi
273	11	1.8	10	22	AAH88269	Hsp-65 peptide epi
274	11	1.8	10	22	AAH88269	Hsp-65 peptide epi
275	11	1.8	15	17	AAH7498	Mycobacterial heat
276	11	1.8	16	17	AAH94826	Peptide from libra
277	11	1.8	16	17	AAH94870	Peptide from libra
278	11	1.8	16	18	AAH43548	Mycobacteria sp. h
279	11	1.8	16	18	AAH43504	Mycobacteria sp. h
280	11	1.8	17	17	AAH94852	Peptide from libra
281	11	1.8	17	18	AAH43530	Mycobacteria sp. h
282	11	1.8	18	22	AAH88292	Hsp-65 peptide epi
283	11	1.8	576	20	AAH23928	Amino acid sequenc
284	11	1.8	577	20	AAH23922	Amino acid sequenc
285	10	1.8	577	20	AAH23927	Amino acid sequenc
286	10	1.8	587	20	AAH23929	Amino acid sequenc
287	9	1.7	9	22	AAH59053	Bacterial conserve
288	9	1.7	9	22	AAH88229	Hsp-65 peptide epi
289	9	1.7	9	22	AAH88230	Hsp-65 peptide epi
290	9	1.7	9	22	AAH88235	Hsp-65 peptide epi
291	9	1.7	9	22	AAH88279	Hsp-65 peptide epi
292	9	1.7	15	16	AAH5802	Immunogenic peptid
293	9	1.7	15	16	AAH81615	Heat shock protein
294	9	1.7	17	17	AAH94783	Peptide from libra
295	9	1.7	17	17	AAH94843	Peptide from libra
296	9	1.7	17	18	AAH43521	Mycobacteria sp. h
297	9	1.7	17	18	AAH43461	Mycobacteria sp. h
298	9	1.7	18	17	AAH94790	Peptide from libra
299	9	1.7	18	18	AAH43468	Mycobacteria sp. h
300	9	1.7	19	17	AAH94854	Peptide from libra
301	9	1.7	19	18	AAH43454	Mycobacteria sp. h
302	9	1.7	19	18	AAH43532	Mycobacteria sp. h
	9	1.7	20	18	AAH33044	Human heat shock p

303	9	1.7	20	18	AAH12352	Human hsp60 peptid
304	9	1.7	114	11	AAH07693	Mycobacterial 65KD
305	9	1.7	558	12	AAH70885	Drosophila melanog
306	8	1.5	9	22	AAH88282	Hsp-65 peptide epi
307	8	1.5	9	22	AAH19467	Mycobacterium sp.
308	8	1.5	9	23	AAH19468	Mycobacterium sp.
309	8	1.5	13	19	AAH54696	Peptide from hsp65
310	8	1.5	15	16	AAH81623	Heat shock protein
311	8	1.5	15	17	AAH81621	Heat shock protein
312	8	1.5	15	17	AAH94847	Peptide from libra
313	8	1.5	15	18	AAH94874	Peptide from libra
314	8	1.5	15	18	AAH43552	Mycobacteria sp. h
315	8	1.5	15	18	AAH43525	Mycobacteria sp. h
316	8	1.5	15	18	AAH19455	Mycobacteria sp. h
317	8	1.5	16	17	AAH94875	Peptide from libra
318	8	1.5	16	17	AAH43553	Mycobacteria sp.
319	8	1.5	16	22	AAH88293	Peptide from libra
320	8	1.5	19	17	AAH94784	Hsp-65 peptide epi
321	8	1.5	19	18	AAH43462	Peptide from libra
322	8	1.5	26	22	AAH88309	Mycobacteria sp. h
323	8	1.5	72	21	AAH33002	Hsp-65 peptide epi
324	8	1.5	93	23	AAH07535	Human OREX protein
325	8	1.5	122	22	AAH03141	Human musculoskele
326	8	1.5	134	21	AAH33001	Arabidopsis thalia
327	8	1.5	156	20	AAH97723	Staphylococcus aur
328	8	1.5	244	21	AAH35964	Zea mays protein f
329	8	1.5	246	20	AAH60028	Human endometrium
330	8	1.5	246	20	AAH60028	Human endometrium
331	8	1.5	256	22	AAH45922	Protonlbacterium
332	8	1.5	287	22	AAH67775	Protonlbacterium
333	8	1.5	338	21	AAH35963	Zea mays protein f
334	8	1.5	364	21	AAH35962	Zea mays protein f
335	8	1.5	440	21	AAH00061	Herbicidially activ
336	8	1.5	524	23	AAH91959	VR22 polypeptide.
337	8	1.5	533	20	AAH85988	Amino acid sequenc
338	8	1.5	545	20	AAH67800	Thermococcus sp. K
339	8	1.5	546	18	AAH26335	KOD-1 heat shock p
340	8	1.5	546	20	AAH67798	Thermococcus sp. K
341	8	1.5	548	20	AAH67797	Thermococcus sp. K
342	8	1.5	549	20	AAH67799	Thermococcus sp. K
343	8	1.5	552	22	AAH67798	Thermococcus sp. K
344	8	1.5	1106	21	AAH03552	Human protein kina
345	8	1.5	1191	21	AAH46051	Arabidopsis thalia
346	8	1.5	1195	21	AAH46050	Arabidopsis thalia
347	8	1.5	1325	21	AAH46049	Novel human diagn
348	8	1.5	1422	22	AAH60345	Novel human diagn
349	8	1.5	1504	22	AAH60358	Drosophila melanog
350	8	1.5	1975	22	AAH62094	Drosophila melanog

ALIGNMENTS

RESULT 1					
ID	AAH23904	AAH23904 standard; Protein; 545 AA.			
AC	AAH23904;				
DT	22-SEP-1999	(first entry)			
DE	Streptococcus pyogenes heat shock protein (Hsp)60-2.				
KW	Heat shock protein; Hsp60-2; Immune response; Immunological carrier;				
OS	Cancer control; Tumour; Sarcoma; Cancer; gene therapy.				
PN	Streptococcus pyogenes.				
PD	MO9935270-A1.				
PF	15-JUL-1999.				
	29-DEC-1998;	98MO-CA01203.			

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XX 31-DEC-1997; 97US-0001737.
PR (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.
XX
XX Mizzen L, Wisniewski J;
XX WPI: 1999-430397/36.
XX N-PSDB: AAX86155.
XX
XX New nucleic acid encoding heat shock protein-60 from Streptococcus,
XX useful in vaccines, as carriers for other immunogens, as anticancer
XX agents and for diagnosis
XX
XX Claim 11: Fig 4A-B; 176pp; English.
XX
XX The present sequence represents a heat shock protein, designated Hsp60-2.
XX The protein, its fragments, variants and fusion proteins, are
XX used to elicit or enhance an immune response against Streptococcus,
XX and to elicit a similar response to a target antigen fused to the
XX protein. Unlike other immunological carriers, Hsp60 proteins are not
XX immunosuppressive so provide an increased response to any conjugated or
XX fused antigen. Also, where used for cancer control, they lack the side
XX effects associated with endotoxins. They can also be used to detect
XX specific antibodies and in treatment or prevention of tumours
XX (e.g. sarcoma or cancers of breast, ovary, prostate, lung, pancreas or
XX liver). The Hsp60 polynucleotide is used for recombinant production
XX of the protein, as a source of primers and probes for detecting
XX streptococci in standard hybridization/amplification assays, and
XX therapeutically in gene therapy vectors.
XX
XX Sequence 545 AA:
XX
XX Query Match 100.0%; Score 545; DB 20; Length 545;
XX Best Local Similarity 100.0%; Pred. No. 0;
XX Matches 545; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 MAKEKFSADARAARVGVMLADTVKVTGKGRNVVLEKAFSPPLITNDGTYTAKKEIE 60
XX 1 MAKEKFSADARAARVGVMLADTVKVTGKGRNVVLEKAFSPPLITNDGTYTAKKEIE 60
XX
XX 61 LEDHFNNGAKLVSEVASKTNDIAGDGTATVLTQALVHEGLKNYTAGANPIGIRGIE 120
XX 61 LEDHFNNGAKLVSEVASKTNDIAGDGTATVLTQALVHEGLKNYTAGANPIGIRGIE 120
XX
XX 121 TATATAVEALKAIAQPVSGKEAIAQVAASVSRSEKVEYISEAMERNGNDGYTTESRG 180
XX 121 TATATAVEALKAIAQPVSGKEAIAQVAASVSRSEKVEYISEAMERNGNDGYTTESRG 180
XX
XX 181 METELEVEEGMOPDRGTLQYWTNENKVAADLENPILITDKKVSNIODILPLEEVLK 240
XX 181 METELEVEEGMOPDRGTLQYWTNENKVAADLENPILITDKKVSNIODILPLEEVLK 240
XX
XX 241 TNRPLLIADVDGEALPTLVNKRGTFFNVVAVKAFGDRRKALMEDIALITGTYIT 300
XX 241 TNRPLLIADVDGEALPTLVNKRGTFFNVVAVKAFGDRRKALMEDIALITGTYIT 300
XX
XX 301 EDLGELEKDATMTALGOAKITVDSDSVYIEGSGSEALANIALIKOLETTTSDPDR 360
XX 301 EDLGELEKDATMTALGOAKITVDSDSVYIEGSGSEALANIALIKOLETTTSDPDR 360
XX
XX 361 EDLGELEKDATMTALGOAKITVDSDSVYIEGSGSEALANIALIKOLETTTSDPDR 360
XX 361 EDLGELEKDATMTALGOAKITVDSDSVYIEGSGSEALANIALIKOLETTTSDPDR 360
XX
XX 421 IEKVALELEGGDAGRNVVLALPEEPRQIALNAGEGSVYIDKLKSPAGTGNATG 480
XX 421 IEKVALELEGGDAGRNVVLALPEEPRQIALNAGEGSVYIDKLKSPAGTGNATG 480
XX
XX 481 EMDVMITGTGIIDPVKVTASALONAAVASLITTEAVVANKPEPATPAPAPAGMDGGM 540
XX 481 EMDVMITGTGIIDPVKVTASALONAAVASLITTEAVVANKPEPATPAPAPAGMDGGM 540
XX
XX DB 481 EMDVMITGTGIIDPVKVTASALONAAVASLITTEAVVANKPEPATPAPAPAGMDGGM 540

```

```

OY 541 GGMG 545
DB 541 GGMG 545
XX
XX RESULT 2
XX ABP28529
XX ID ABP28529 standard; Protein: 545 AA.
XX
XX ABP28529;
XX
XX 02-JUL-2002 (first entry)
XX
XX Streptococcus polypeptide SEQ ID NO 6234.
XX
XX Streptococcus; GAS; GBS; group B streptococcus; Streptococcus agalactiae;
XX group A streptococcus; Streptococcus pyogenes; antibacterial;
XX antiinflammatory; infection; vaccine; meningitis; gene therapy.
XX
XX Streptococcus pyogenes.
XX
XX MO200234771-A2.
XX
XX 02-MAY-2002.
XX
XX 29-OCT-2001; 2001MO-GB04789.
XX
XX 27-OCT-2000; 2000GB-0026333.
XX 24-NOV-2000; 2000GB-0028727.
XX 07-MAR-2001; 2001GB-0005640.
XX
XX (CHIR-) CHIRON SPA.
XX (GENO-) INST GENOMIC RES.
XX
XX Telford J, Maignan V, Margalit Ros YI, Grandi G, Fraser C;
XX Telford J;
XX
XX WPI: 2002-35236/38.
XX N-PSDB: ABN69160.
XX
XX New Streptococcus protein for the treatment or prevention of infection
XX or disease caused by Streptococcus bacteria, such as meningitis, and
XX for detecting a compound that binds to the protein -
XX
XX Claim 1: Page 3786; 4525pp; English.
XX
XX The invention relates to a protein (ABP25413-ABP30895) from group B
XX streptococcus/GBS (Streptococcus agalactiae) or group A streptococcus/GAS
XX (Streptococcus pyogenes), comprising one of 5483 sequences (S1), given in
XX the specification. The proteins have antibacterial and antiinflammatory
XX activity. (I) nucleic acids encoding (I), ABN6044-ABN71526 and
XX antibodies that bind (I) are used in the manufacture of medicaments for
XX the treatment or prevention of infection or disease caused by
XX Streptococcus bacteria, particularly S. agalactiae and S. pyogenes.
XX Nucleic acids encoding (I) are used to detect Streptococcus in a
XX biological sample. (I) is used to determine whether a compound binds to
XX (I). A composition comprising (I) or a nucleic acid encoding (I), may be
XX used as a vaccine or diagnostic composition. The disease caused by
XX Streptococcus that is prevented or treated may be meningitis. Nucleic
XX acid encoding (I) may be used to recombinantly produce (I) and may be
XX used in gene therapy. Antibodies to (I) are used for affinity
XX chromatography, immunoassays, and distinguishing/identifying
XX Streptococcus proteins.
XX
XX Sequence 545 AA:
XX
XX Query Match 98.7%; Score 538; DB 23; Length 545;
XX Best Local Similarity 100.0%; Pred. No. 0;
XX Matches 538; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 5 IKFSADARAARVGVMLADTVKVTGKGRNVVLEKAFSPPLITNDGTYTAKKEIE 64
XX 7 IKFSADARAARVGVMLADTVKVTGKGRNVVLEKAFSPPLITNDGTYTAKKEIE 64
XX
XX DB 7 IKFSADARAARVGVMLADTVKVTGKGRNVVLEKAFSPPLITNDGTYTAKKEIE 64

```

QY 65 FENWGAIVSEVASKNDIAGDCTTATVLTQAIHGEGLKNVAGANPIGIRGIEPTATA 124
 DB 67 FENWGAIVSEVASKNDIAGDCTTATVLTQAIHGEGLKNVAGANPIGIRGIEPTATA 126
 QY 125 TAVEALKAIAPVSGKEAIAQVAAVSSREKVEYISEAMERVNDGVITIEESRGME 184
 DB 127 TAVEALKAIAPVSGKEAIAQVAAVSSREKVEYISEAMERVNDGVITIEESRGME 186
 QY 185 LEVEGEGPDRGLISQYMTDNEKRVADLENPFLITDKVSNIOIILPLEEVLTNRP 244
 DB 187 LEVEGEGPDRGLISQYMTDNEKRVADLENPFLITDKVSNIOIILPLEEVLTNRP 246
 QY 245 LLIIADVDGEALPVLVNLKRGTFENNVAVAPGFGDRRAKMEIDAILTGCTVTEDLG 304
 DB 247 LLIIADVDGEALPVLVNLKRGTFENNVAVAPGFGDRRAKMEIDAILTGCTVTEDLG 306
 QY 305 LEIKDATMTALGOAKITVDKSTIYVSGSSSAINRIALIKSOLLETTSDPDRKLO 364
 DB 307 LEIKDATMTALGOAKITVDKSTIYVSGSSSAINRIALIKSOLLETTSDPDRKLO 366
 QY 365 ERLAKLAGVAVIKVGAFTETALKEMKLRLEDALNATRAAVEGIVAGGTALITVIEKV 424
 DB 367 ERLAKLAGVAVIKVGAFTETALKEMKLRLEDALNATRAAVEGIVAGGTALITVIEKV 426
 QY 425 AALEEGDATTGRNIVLRALPEPVQIALNAGYEGSVYIDKLNPSAGTGFAATGEVND 484
 DB 427 AALEEGDATTGRNIVLRALPEPVQIALNAGYEGSVYIDKLNPSAGTGFAATGEVND 486
 QY 485 MITGIIIDPVKVTTRSAALNAAVSLITTEAVVANKPEPATAPAMPAGMDPGMGG 542
 DB 487 MITGIIIDPVKVTTRSAALNAAVSLITTEAVVANKPEPATAPAMPAGMDPGMGG 544
 RESULT 3
 ABP28528
 ID ABP28528 standard: Protein: 540. AA.
 AC ABP28528;
 DT 02-JUL-2002 (first entry)
 DE Streptococcus polypeptide SEQ ID NO 6232.
 XX Streptococcus: GAS; group B streptococcus; Streptococcus agalactiae;
 KM group A streptococcus; Streptococcus pyogenes; antibacterial;
 KM antiinflammatory; infection; vaccine; meningitis; gene therapy.
 OS Streptococcus agalactiae.
 XX
 FN WO200234771-A2.
 PD 02-MAY-2002.
 PF 29-OCT-2001; 2001WO-GB04789.
 XX 27-OCT-2000; 2000GB-0026333.
 PR 24-NOV-2000; 2000GB-0028727.
 PR 07-MAR-2001; 2001GB-0005640.
 PA (CHIR-) CHIRON SPA.
 PA (GENO-) INST GENOMIC RES.
 PI Telford J, Masignani V, Margarit Ros YI, Grandi G, Fraser C;
 PI Tettelin H.
 XX WPI: 2002-352536/38.
 DR N-PSDB: ABN69159.
 XX New Streptococcus protein for the treatment or prevention of infection
 PT or disease caused by Streptococcus bacteria, such as meningitis, and
 PT for detecting a compound that binds to the protein -
 XX

PS Claim 1; Page 3785; 4525pp; English.
 XX
 CC The invention relates to a protein (ABP25413-ABP30895) from group B
 CC Streptococcus/GBS (Streptococcus agalactiae) or group A streptococcus/GAS
 CC (Streptococcus pyogenes), comprising one of 5483 sequences (S1), given in
 CC the specification. The proteins have antibacterial and antiinflammatory
 CC activity. (1), nucleic acids encoding (1), ABN6044-ABN71526 and
 CC antibodies that bind (1) are used in the manufacture of medicaments for
 CC the treatment or prevention of infection or disease caused by
 CC Streptococcus bacteria, particularly S. agalactiae and S. pyogenes;
 CC Nucleic acids encoding (1) are used to detect streptococcus in a
 CC biological sample. (1) is used to determine whether a compound binds to
 CC (1). A composition comprising (1) or a nucleic acid encoding (1), may be
 CC used as a vaccine or diagnostic composition. The disease caused by
 CC streptococcus that is prevented or treated may be meningitis. Nucleic
 CC acid encoding (1) may be used to recombinantly produce (1) and may be
 CC used in gene therapy. Antibodies to (1) are used for affinity
 CC chromatography, immunoassays, and distinguishing/identifying
 CC Streptococcus proteins.
 CC
 S0 Sequence 540 AA:
 Query Match 14.3%; Score 78; DB 23; Length 540;
 Best Local Similarity 100.0%; Pred. No. 3e-66;
 Matches 78; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 22 LAOTVAVTLGPRGVNVEKAFGSPLTINDGVITAKEIELEDFENWGAIVSEVASKTN 81
 DB 22 LAOTVAVTLGPRGVNVEKAFGSPLTINDGVITAKEIELEDFENWGAIVSEVASKTN 81
 QY 82 DIAGDCTTATVLTQAI 99
 DB 82 DIAGDCTTATVLTQAI 99
 RESULT 4
 AAM01101
 ID AAM01101 standard: Protein: 540. AA.
 AC AAM01101;
 DT 02-OCT-2001 (first entry)
 DE CFE 104 protein sequence.
 XX
 DE Antibacterial; vaccine; gene therapy; bacterial cell wall viability;
 KM CFE; CEG; Conserved Essential Gene; bacterial infection;
 KM antisense therapy; antibiotic resistance.
 XX
 OS Streptococcus pneumoniae.
 XX
 FN WO200149721-A2.
 PD 12-JUL-2001.
 PF 29-DEC-2000; 2000WO-US35604.
 XX 30-DEC-1999; 99US-0174089.
 PR (BRIM) BRISTOL-MYERS SQUIBB CO.
 PA
 PA
 PI Dougherty TJ, Pucci MD, Dougherty BA, Davison DB, Brucoleri RE;
 PI Thannasi JA.
 XX WPI: 2001-496721/54.
 DR N-PSDB: AAH90800.
 XX Nucleic acids encoding conserved essential genes involved in bacterial
 PT replication which are potential targets for the treatment of antibiotic
 PT resistant bacterial infections -
 XX
 PS Claim 27; Pages 356-358; 380pp; English.

CC The present invention relates to nucleic acids (AAH90701-AAH90918).
 CC encoding polypeptides (AAH01002-AAH0114), which are essential for the
 CC viability of a bacterial cell wall. The acronym CFE stands for "CEG For
 CC Expression", where CEG stands for "Conserved Essential Gene". The nucleic
 CC acids are useful for detecting the presence of proteins essential for the
 CC viability of a bacterial cell wall in samples such as cells, tissues,
 CC biological fluids, blood, serum, nose, ear or throat swabs with ligands,
 CC and for detecting corresponding target nucleic acid molecules with
 CC complementary sequences. The nucleic acids are also useful for
 CC determining whether a genomic nucleotide sequence of interest is
 CC essential for viability of a bacterial cell or whether it resides within
 CC an operon. By integrating an exogenous nucleotide sequence comprising a
 CC portion of an open reading frame of the genomic sequence of interest
 CC (comprising 200-500 base pairs) into the genomic sequence of interest
 CC which confers a selectable phenotype to the cell, and determining cell
 CC viability with a selection agent such as chloramphenicol. The nucleic
 CC acids and proteins are also useful as vaccines and for treating bacterial
 CC infections with gene therapy and antisense therapy. The nucleic acids
 CC also enable identification of targets suitable for the treatment of
 CC antibiotic resistant bacterial infections.

XX Sequence 540 AA;

Query Match 12.8%; Score 70; DB 22; Length 540;

Best Local Similarity 100.0%; Pred. No. 1.6e-58;

Matches 70; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 242 NRPLIIADVDGEALPTLVKIRGTNNVAVKAPGDRRKAMEIDAIITGCTVITE 301
 DB 242 NRPLIIADVDGEALPTLVKIRGTNNVAVKAPGDRRKAMEIDAIITGCTVITE 301
 OY 302 DLGLEKDAT 311
 DB 302 DLGLEKDAT 311

RESULT 5

AAV23902

ID AAV23902 standard; Protein: 541 AA.

XX AC AAV23902;

DT 22-SEP-1999 (first entry)

DE Streptococcus pneumoniae heat shock protein (Hsp)60-2.

KW Heat shock protein; Hsp60-2; immune response; immunological carrier;
 cancer control; tumour; sarcoma; cancer; gene therapy.

OS Streptococcus pneumoniae.

PN MO9935270-A1.

PD 15-JUL-1999.

PF 29-DEC-1998; 98WO-CA01203.

PR 31-DEC-1997; 97US-0001737.

PI (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.

PA Mizzen L, Wisniewski J;

PI Mizzen L, Wisniewski J;

DR WPI; 1999-430397/36.

DR N-PSDB; AAX86153.

PT New nucleic acid encoding heat shock protein-60 from Streptococcus,
 useful in vaccines, as carriers for other immunogens, as anticancer
 agents and for diagnosis.

PS Claim 11; Fig 2A-B; 176pp; English.

CC The present sequence represents a heat shock protein, designated Hsp60-2.

CC The protein, its fragments, variants and fusion proteins, are
 CC used to elicit or enhance an immune response against Streptococcus,
 CC and to elicit a similar response to a target antigen fused to the
 CC protein. Unlike other immunological carriers, Hsp60 proteins are not
 CC immunosuppressive so provide an increased response to any conjugated or
 CC fused antigen. Also, where used for cancer control, they lack the side
 CC effects associated with endotoxins. They can also be used to detect
 CC specific antibodies and in treatment or prevention of tumours
 CC (e.g. sarcoma or cancer of breast, ovary, prostate, lung, pancreas or
 CC liver). The Hsp60 polynucleotide is used for recombinant production
 CC of the protein, as a source of primers and probes for detecting
 CC streptococci in standard hybridization/amplification assays, and
 CC therapeutically in gene therapy vectors.

XX Sequence 541 AA;

Query Match 12.8%; Score 70; DB 20; Length 541;

Best Local Similarity 100.0%; Pred. No. 1.6e-58;

Matches 70; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 242 NRPLIIADVDGEALPTLVKIRGTNNVAVKAPGDRRKAMEIDAIITGCTVITE 301
 DB 242 NRPLIIADVDGEALPTLVKIRGTNNVAVKAPGDRRKAMEIDAIITGCTVITE 301
 OY 302 DLGLEKDAT 311
 DB 302 DLGLEKDAT 311

RESULT 6

AAH31619

ID AAB31619 standard; Protein: 641 AA.

XX AC AAB31619;

DT 30-APR-2001 (first entry)

DE Amino acid sequence of Hsp65-E7 fusion protein.

KW Heat shock protein; Hsp; Th1 response; Th1 cell; CD4+ T lymphocyte cell;
 lymphocyte; Hsp65; Hsp60; Hsp10; Hsp60; Hsp71; microbial pathogen;
 E7 protein.

OS Synthetic.

OS Streptococcus pneumoniae.

OS Human papillomavirus.

PN WO200104344-A2.

PD 18-JAN-2001.

PF 10-JUL-2000; 2000WO-US18828.

PR 08-JUL-1999; 99US-0143757.

PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.

PI Siegel M, Chu NR, Mizzen LA;

DR WPI; 2001-138361/14.

DR N-PSDB; AAF25036.

PT Screening for compounds that stimulate Th1-like responses in CD4+ T
 lymphocyte cells

PS Example 15; Fig 15A-B; 88pp; English.

CC The present sequence represents a fusion protein comprising a
 CC Streptococcus pneumoniae heat shock protein (Hsp) 65 fused to a HPV16 E7
 CC protein. The fusion protein is used in the method of the invention. The
 CC specification describes a method of determining whether a compound
 CC stimulates a Th1-like response. Th1 cells are a subset of CD4+
 CC T lymphocyte cells. The method comprises contacting naive lymphocytes

CC In vitro with a fusion protein comprising at least a fragment of Hsp.
CC and then detecting the Th1-like response exhibited by the cell sample.
CC The proteins which may be used in the method of the invention are Hsp65,
CC Hsp40, Hsp10, Hsp60, and Hsp71. The method may be used to identify
CC compounds that stimulate Th1-like responses in response to microbial
CC pathogens.
XX

SO Sequence 641 AA;

Query Match 12.8%; Score 70; DB 22; Length 641;
Best Local Similarity 100.0%; Pred. No. 1.9e-58;
Matches 70; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 242 NRPLIADVDGALPTLVNKRGTENVAVKAPGFCRRKAMLEDAITLGTGYTE 301
DB 242 NRPLIADVDGALPTLVNKRGTENVAVKAPGFCRRKAMLEDAITLGTGYTE 301
OY 302 DGLLEKDAT 311
DB 302 DGLLEKDAT 311

RESULT 7
ID ABB53701 standard; Protein; 542 AA.

AC ABB53701;
XX 16-MAY-2002 (first entry)
XX Lactococcus lactis protein groEL.
XX Lactococcus lactis protein groEL.
XX Biosynthesis; biodegradation; lactic bacterium; yogurt; cheese.
XX Lactococcus lactis IL1403.
XX FR2807446-A1.
XX 12-OCT-2001.

XX 11-APR-2000; 2000FR-0004630.
XX 11-APR-2000; 2000FR-0004630.
XX (INRG) INRA INST NAT RECH AGRONOMIQUE.

PI Bolocline A, Sorokline A, Renault P, Ehrlich SD;
XX WPI; 2002-043418/06.

PT New nucleotide sequence useful in the identification of Lactococcus
XX lactis and related species.

PS Claim 6; SEQ ID No 403; 2504pp; French.

CC The present invention is related to a Lactococcus lactis nucleotide
CC sequence (AB590521) and related proteins (AB53300-AB55621). The
CC nucleic acid sequence is useful in the detection and/or amplification of
CC related species. The proteins of the invention are useful for the
CC biosynthesis or biodegradation of a composition of interest. The
CC invention helps research in lactic bacteria, particularly useful in the
CC production of yogurt and cheese.
CC Note: The sequence data for this patent is based on equivalent patent
CC WO200177334 (published 18-OCT-2001) which is available in electronic
CC format directly from WIPO at ftp.wipo.int/pub/published_pcl_sequences.
XX

SO Sequence 542 AA;

Query Match 10.3%; Score 56; DB 23; Length 542;
Best Local Similarity 100.0%; Pred. No. 5.1e-45;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 44 GSPILINDGVTIANKIELEDFHFMGAKLVSEVASKNDIAGDGTATVLTQAIY 99
DB 44 GSPILINDGVTIANKIELEDFHFMGAKLVSEVASKNDIAGDGTATVLTQAIY 99

RESULT 8

ID ABB49241 standard; Protein; 542 AA.

AC ABB49241;
XX 05-FEB-2002 (first entry)

XX Listeria monocytogenes protein #1945.
XX Antibacterial; gene therapy; vaccine; biosynthesis; biodegradation;
XX vitamin B12; bacterial infection; disease.
XX Listeria monocytogenes.

XX WO200177335-A2.

XX 18-OCT-2001.
XX 11-APR-2001; 2001NO-FR01118.
XX 11-APR-2000; 2000FR-0004629.

XX (INSP) INST PASTEUR.
XX Buchrieser C, Franquel L, Couve E, Rusniok C, Fsihi H, Dehoux P;
XX Dussurget O, Chetoui M, Nedjari H, Glaser P, Kunst F, Cossart P;
XX Daniels J, Goebel W, Kreft J, Kuhn M, Ng E, Vazquez-Boland JA;
XX Dominguez-Bernal G, Garrido-Garcia P, Tierrez-Martinez A, Amend A;
XX Chakraborty T, Dommann E, Hahn T, Berche P, Chardot A, Durant L;
XX Perez-Diaz J, Baquero F, Garcia Del Portillo F, Gomez-Lopez N;
XX Madueno E, De Pablos B, Wehlund J, Kaerst U, Entian K, Hauf J;
XX Rose M, Voss H;

XX WPI; 2002-010914/01.
XX Genomic sequence for Listeria monocytogenes, useful e.g. for treatment
XX and prevention of Listeria and related bacterial infections, and
XX related polypeptides

PT Claim 6; SEQ ID No 1946; 192pp; French.
XX The present invention relates to the genome sequence of Listeria
XX monocytogenes EGD-e (see ABA03041). The genome sequence and fragments of
XX it are useful for selecting probes and primers for detecting genes in L.
XX monocytogenes and related organisms, and for studying genetic
XX polymorphisms and other genomes. The present sequence is a protein
XX encoded by the genome sequence of the present invention. Proteins
XX expressed from the genome sequence are useful for raising specific
XX antibodies, identification of L. monocytogenes and related organisms, and
XX for biosynthesis and biodegradation, especially biosynthesis of vitamin
XX B12. The genome sequence and proteins encoded by it are also useful for
XX selecting compounds that regulate gene expression and cell replication
XX and modulate L. monocytogenes-related diseases. In addition, the genome
XX sequence and proteins encoded by it are useful in pharmaceutical and
XX vaccines compositions for the treatment or prevention of infections by L.
XX monocytogenes and related organisms.

CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX

SO Sequence 542 AA;

Query Match 5.7%; Score 31; DB 23; Length 542;
Best Local Similarity 100.0%; Pred. No. 6.6e-21;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 266 RGTENVAVKAPGFCRRKAMLEDAITLGTG 296

XX	Amino acid sequence of a Chlamydia trachomatis protein.
DE	Vaccine; eye disease; conventional trachoma; nongendemic trachoma;
KW	paratrachoma; inclusion conjunctivitis; genital diseases; perithelatitis;
KW	nongonococcal urethritis; epididymitis; cervicitis; salpingitis;
KV	bartholinits; pneumonia; venereal lymphogranulomatosis.
XX	Chlamydia trachomatis.
OS	M09928475-A2.
PN	10-JUN-1999..
PD	27-NOV-1998: 98WO-IB01939.
XX	04-NOV-1998: 98US-0107077.
XX	28-NOV-1997: 97ER-0015041.
PR	17-DEC-1997: 97ER-0016034.
PA	(GEST) GENSET.
PX	Grieffals R:
PI	WPI; 1999-371125/31.
DR	Genome sequence of Chlamydia trachomatis
XX	Disclosure: Page 900: 1755pp: English.
PS	AAV36754-Y37949 are encoded by open reading frames (ORFs) of the genome
CC	of Chlamydia trachomatis (see AF01425). The polypeptides can be used as
CC	vaccines against Chlamydia trachomatis. Antisense and ribozyme sequences
CC	can also be used to control growth of the microorganism. Chlamydia
CC	trachomatis is responsible for a large number of diseases, e.g. eye
CC	diseases such as conventional trachoma, nongendemic trachoma,
CC	paratrachoma, and inclusion conjunctivitis; genital diseases such as
CC	nongonococcal urethritis, epididymitis, cervicitis, salpingitis,
CC	perithelitis, bartholinits; pneumonia; venereal lymphogranulomatosis.
CC	The polypeptides of the invention may be of use in treating these diseases.
CC	Sequence 224 AA:
SQ	
Query Match	4.4% Score 24: DB 20: Length 224:
Best Local Similarity	100.0%; Pred. No. 1,6e-14:
Matches 24: Conservative	0: Mismatches 0: Indels 0: Gaps 0:
OY	273 AVKAPGFGRRAKMEIDAILTGG 296
DB	20 AVKAPGFGRRAKMEIDAILTGG 43
RESULT 11	
AAAI0977	ID AAMI0977 standard: Protein: 419 AA.
AC	AAAI0977:
XX	21-MAY-1997 (first entry)
DE	Dihydrofolate reductase-Chlamydia pneumoniae antigen fusion protein.
XX	DHFR, dihydrofolate reductase: Chlamydia pneumoniae; pneumonia;
KM	antibody production; diagnosis; fusion protein.
XX	Chlamydia pneumoniae (chimeric).
OS	Location/Qualifiers
FH	Key
FT	Region
FT	161..170
FT	/note-"peptide linker"
TT	/note-"dihydrofolate reductase region"

FT Misc-difference 171..407
 FT /note= "Chlamydia pneumoniae antigen region"
 XX JP08294391-A.
 XX
 PD 12-NOV-1996.
 XX
 XX 28-APR-1995; 95JP-0106007.
 XX
 XX 28-APR-1995; 95JP-0106007.
 XX
 PA (HITB) HITACHI CHEM CO LTD.
 XX
 DR WPI; 1997-036901/04.
 XX
 PT Fusion protein comprising dihydrofolate reductase and Chlamydia
 PT pneumoniae antigen - useful in prodn. of C. pneumoniae antibodies
 PT for diagnosis of infection
 XX
 PS Claim 5; Page 14-15; 17pp; Japanese.
 XX
 CC AAM10977 is a dihydrofolate reductase (DHFR)-Chlamydia pneumoniae
 CC antigen fusion protein. Fusion proteins that may be made consist of at
 CC least 5 contiguous amino acids of the Chlamydia pneumoniae antigen
 CC linked to the N-terminus of a DHFR protein (see AAM10974). Fusion
 CC proteins produced are useful for the production of anti-C. pneumoniae
 CC antibodies which are useful in the diagnosis and treatment of infectious
 CC diseases caused by C. pneumoniae.
 XX
 SQ Sequence 419 AA;
 XX
 Query Match 4.4%; Score 24; DB 18; Length 419;
 Best Local Similarity 100.0%; Pred. No. 2.9e-14;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 273 AVKAPGFGRRKAMEDIAITLGG 296
 DB 243 AVKAPGFGRRKAMEDIAITLGG 266
 XX
 RESULT 12
 AAM1865
 ID AAM1865 standard; Protein; 419 AA.
 XX
 AC AAM1865;
 XX
 XX 17-APR-1997 (first entry)
 DE DHFR/Polypeptide B fusion protein #2.
 XX
 KM Dihydrofolate reductase; DHFR; C. pneumoniae; detection; antibody;
 KM fusion protein; antigen; diagnosis.
 XX
 OS Chlamydia pneumoniae.
 XX
 FH Key Location/Qualifiers
 FT Protein 1..160
 FT /label= DHFR
 FT 172..407
 FT /label= Polypeptide_B_residues_203-439
 XX
 PN JP08304403-A.
 XX
 XX 22-NOV-1996.
 XX
 XX 28-APR-1995; 95JP-0106012.
 XX
 XX 28-APR-1995; 95JP-0106012.
 XX
 PA (HITB) HITACHI CHEM CO LTD.
 XX
 DR WPI; 1997-056177/06.
 XX

PT Detection and determination of anti-Chlamydia pneumoniae antibody -
 PT using an antigenic polypeptide fused to the Chlamydia
 PT dihydrofolate reductase as the antigen
 XX
 PS Claim 5; Page 15-16; 17pp; Japanese.
 XX
 CC This sequence represents a fusion between C. pneumoniae polypeptide
 CC B and dihydrofolate reductase (DHFR). This protein was used in the
 CC method of the invention for the detection and determination of
 CC anti-Chlamydia pneumoniae antibody. DHFR and polypeptide B were
 CC combined directly or through an amino acid sequence to give a fusion
 CC protein to act as an antigen. The fusion protein may be used in
 CC a reagent which has a high reliability and which gives an exact
 CC diagnosis.
 XX
 SQ Sequence 419 AA;
 XX
 Query Match 4.4%; Score 24; DB 18; Length 419;
 Best Local Similarity 100.0%; Pred. No. 2.9e-14;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 273 AVKAPGFGRRKAMEDIAITLGG 296
 DB 243 AVKAPGFGRRKAMEDIAITLGG 266
 XX
 RESULT 13
 AAR13337
 ID AAR13337 standard; Protein; 544 AA.
 XX
 AC AAR13337;
 XX
 DT 17-DEC-2001 (updated)
 DT 22-OCT-1991 (first entry)
 XX
 DE Hyb protein.
 XX
 KM Antibodies; heat shock; hypersensitive; allergen; HSP60; GROES;
 KM GREL.
 XX
 OS Chlamydia trachomatis serovar A.
 XX
 PN USN/531317-N.
 XX
 PD 09-JUL-1991.
 XX
 PF 31-MAY-1990; 90US-0143560.
 XX
 PR 31-MAY-1990; 90US-0531317.
 XX
 XX (USSH) NAT INST OF HEALTH.
 PA WPI; 1991-245693/33.
 XX
 DR N-PSDB; AAQ13137.
 XX
 XX DNA encoding HybA and HybB Chlamydia proteins - used to develop
 PT prods. for detection of and vaccines against Chlamydia infection.
 PT
 XX
 PS Disclosure; Fig 7; 51pp; English.
 XX
 CC The sequence was deduced from the second of two ORFs found in the
 CC hyp operon of clone pTAS71CC prepd. from C. trachomatis genomic DNA.
 CC It is the HybB hypersensitivity protein, analogous to the E. coli
 CC GREL protein. It can be used to raise antibodies and to
 CC prepare vaccines for the treatment of Chlamydial infections. The
 CC protein also elicits a cell-mediated immune response so can be used
 CC as a skin test antigen.
 CC See also AAR13334-R13336.
 CC (Note: Revised entry submitted to correct the patent number format
 CC US Government-owned NIS applications to prevent classes with ongc
 CC granted patent numbers. For further information please visit the
 CC web site at www.derwent.com/dwpi/updates/ntis_us.html.)
 XX

SO Sequence 544 AA:

Query Match 4.4%; Score 24; DB 12; Length 544;
Best Local Similarity 100.0%; Pred. No. 3.7e-14;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 273 AVKAPGFGDRRKAMLEDAITLTGG 296
|||||
DB 275 AVKAPGFGDRRKAMLEDAITLTGG 298

RESULT 14

AA67383 standard; Protein; 544 AA.

AA67383;

22-JUN-1995 (first entry)

C. psittaci Hyps gene product.

Urease; immunogen; vaccine; diagnostic; heat shock protein; HSP;

GroEL-like protein; Helicobacter felis.

Chlamydia psittaci.

MO9426901-A.

24-NOV-1994.

19-MAY-1994; 94MO-EP01625.

19-MAY-1993; 93EP-0401309.

19-NOV-1993; 93WO-EP03259.

(INRM) INST NAT SANTE & RECH MEDICALE.

(INSP) INST PASTEUR.

Ferrero R, Labigne A, Suerbaum S, Thiberge J;

WPI; 1995-006797/01.

DNA from Helicobacter pylori and Helicobacter felis - used to

develop prods. for detection, treatment and prevention of

Helicobacter infection

Disclosure: Fig. 7A(i-vii); 168pp; English.

The sequence of the Helicobacter pylori heat shock protein A

(given in AA67374) was compared to that of other GroEL-like

proteins from Legionella pneumophila (AA67381), Escherichia coli

(AA67382), Chlamydia psittaci (AA67383), Mycobacterium leprae

(AA67384) and human mitochondrial protein PI (AA67385), and regions

of homology were identified.

Sequence 544 AA;

Query Match

Best Local Similarity 100.0%; Score 24; DB 16; Length 544;

Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 273 AVKAPGFGDRRKAMLEDAITLTGG 296
|||||
DB 275 AVKAPGFGDRRKAMLEDAITLTGG 298

RESULT 15

AA67383 standard; Protein; 544 AA.

AA67383;

21-MAY-1997 (first entry)

Chlamydia pneumoniae antigen used as DHFR-linked fusion protein.
DHFR: dihydrofolate reductase; Chlamydia pneumoniae; pneumoniae;
antibody production; diagnosis; fusion protein.

Chlamydia pneumoniae.

Key Location/Qualifiers
Misc-difference 25 /note- "given as Gie in three letter amino
acid code in the specification"

JP08294391-A.

12-NOV-1996.

28-APR-1995; 95JP-0106007.

28-APR-1995; 95JP-0106007.

(HITB) HITACHI CHEM CO LTD.

WPI; 1997-036901/04.

Fusion protein comprising dihydrofolate reductase and Chlamydia
pneumoniae antigen - useful in prodn. of C. pneumoniae antibodies
for diagnosis of infection

Claim 1; Page 11-12; 17pp; Japanese.

AA67383 encodes a 544 residue Chlamydia pneumoniae antigen, at least
5 contiguous amino acids of which are fused to a dihydrofolate
reductase (DHFR) enzyme. Fusion proteins produced are useful for the
production of anti-C. pneumoniae antibodies which are useful in the
diagnosis and treatment of infectious diseases caused by C. pneumoniae.

Sequence 544 AA;

Query Match 4.4%; Score 24; DB 18; Length 544;

Best Local Similarity 100.0%; Pred. No. 3.7e-14;

Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 273 AVKAPGFGDRRKAMLEDAITLTGG 296
|||||
DB 275 AVKAPGFGDRRKAMLEDAITLTGG 298

Search completed: April 8, 2003, 14:35:09
Job time : 81 secs